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APPLICATION NO. FILING DATE		FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
09/900,115	07/06/2001	Stephen H. Bartelmez	50450-8039.US00	50450-8039.US00 5757	
22918	7590 09 19.2002				
PERKINS COIE LLP			EXAMINER		
P.O. BOX 216 MENLO PAR	68 .K, CA 94026	TON, THAIAN N			
			ART UNIT	PAPER NUMBER	
			1632	K	
			DATE MAILED: 09.19.2002	1)	

Please find below and/or attached an Office communication concerning this application or proceeding.

,		Application	n No.	Applicant(s)				
Office Assistant Communication		09/900,115	5	BARTELMEZ ET AL.				
	Office Action Summary	Examin r		Art Unit				
		Thaian N.		1632				
The MAILING DATE of this communication appears on the cover sheet with the correspondence address Period for Reply								
A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 1 MONTH(S) FROM THE MAILING DATE OF THIS COMMUNICATION. - Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication. - If the period for reply specified above is less than thirty (30) days, a reply within the statutory minimum of thirty (30) days will be considered timely. - If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication. - Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). - Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b). Status								
1)	Responsive to communication(s) filed on							
2a)	This action is FINAL 2b) Th	is action is i	non-final.					
3) Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under <i>Ex parte Quayle</i> , 1935 C.D. 11, 453 O.G. 213.								
Disposition of Claims								
4) Claim(s) 1-21 is/are pending in the application.								
4a) Of the above claim(s) is/are withdrawn from consideration.								
5) Claim(s) is/are allowed.								
6) Claim(s) is/are rejected.								
7)	7) Claim(s) is/are objected to.							
8) Claim(s) 1-21 are subject to restriction and/or election requirement.								
Applicati	on Papers							
, —	The specification is objected to by the Examine							
10) The drawing(s) filed on is/are: a) accepted or b) objected to by the Examiner.								
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).								
11) The proposed drawing correction filed on is: a) □ approved b) □ disapproved by the Examiner.								
If approved, corrected drawings are required in reply to this Office action.								
12) The oath or declaration is objected to by the Examiner.								
Priority under 35 U.S.C. §§ 119 and 120								
13) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).								
a) All b) Some * c) None of:								
1. Certified copies of the priority documents have been received.								
	2. Certified copies of the priority documents have been received in Application No							
 3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)). * See the attached detailed Office action for a list of the certified copies not received. 								
14) Acknowledgment is made of a claim for domestic priority under 35 U.S.C. § 119(e) (to a provisional application).								
a) ☐ The translation of the foreign language provisional application has been received. 15) ☐ Acknowledgment is made of a claim for domestic priority under 35 U.S.C. §§ 120 and/or 121.								
Attachment(s)								
1) Notic	te of References Cited (PTO-892) se of Draftsperson's Patent Drawing Review (PTO-948) mation Disclosure Statement(s) (PTO-1449) Paper No(s) _	·		y (PTO-413) Paper No Patent Application (P				

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DETAILED ACTION

Election/Restrictions

Restriction to one of the following inventions is required under 35 U.S.C. 121:

- I. Claims 1·14 drawn to methods of prolonging the survival time of human stem cells in culture by exposing an enriched stem cell population to an oligomer antisense to TGF·β, and a composition of human stem cells treated with an oligomer antisense to TGF·β, classified in class 435, subclasses 325, 363, 366, 377, 404, for example.
- II. Claims 15-18, 20 and 21, drawn to methods of decreasing the time for hematopoietic reconstitution of a patient following chemotherapy or radiation therapy, classified in class 514, subclass 44, class 325, 363, 366, 377, for example.
- III. Claim 19, drawn to a method of rapid *in vitro* production of lineage committed progenitor cells and their progeny from human stem cells, classified in class 435, subclasses 325, 364, 366, 377, for example.

The inventions are distinct, each from the other because of the following reasons:

Inventions I and either of Inventions II or III are related as product and process of use. The inventions can be shown to be distinct if either or both of the following can be shown: (1) the process for using the product as claimed can be practiced with another materially different product or (2) the product as claimed can be used in a materially different process of using that product (MPEP § 806.05(h)). In the instant case the human stem cell composition of Invention I can be used as a source of undifferentiated stem cells.

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Inventions II and III are mutually exclusive and independent. The methods of decreasing the time for hematopoietic reconstitution of a patient following chemotherapy or radiation therapy of Invention II are not required for the implementation of the method of rapid *in vitro* of lineage committed progenitor cells and their progeny from human stem cells of Invention III, and vice versa. Furthermore, each of the methods requires a materially different and separate protocol.

Because these inventions are distinct for the reasons given above and have acquired a separate status in the art because of their recognized divergent subject matter, restriction for examination purposes as indicated is proper.

Applicant is advised that the reply to this requirement to be complete must include an election of the invention to be examined even though the requirement be traversed (37 CFR 1.143).

Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a request under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

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Sequence Election Requirement

Claims 1-50 read on patentably distinct Groups drawn to multiple SEQ ID Numbers [in particular, SEQ ID NO: 1, 2 or 5]. Each of the SEQ ID Nos constitutes independent inventions, which are patentably distinct because each of the sequences are unrelated, as such a further restriction is applied to the sequences. For an elected Group drawn to a SEQ ID NO., the Applicants must further elect a single sequence.

MPEP 803.04 states:

Nucleotide sequences encoding different proteins are structurally distinct chemical compounds and are unrelated to one another. These sequences are thus deemed to normally constitute independent and distinct inventions within the meaning of 35 U.S.C. 121. Absent evidence to the contrary, each such nucleotide sequence is presumed to represent an independent and distinct invention, subject to a restriction requirement pursuant to 35 U.S.C. 121 and 37 CFR 1.141 et seq. Nevertheless, to further aid the biotechnology industry in protecting its intellectual property without creating an undue burden on the Office, the Commissioner has decided sua sponte to partially waive the requirements of 37 CFR 1.141 et seq. and permit a reasonable number of such nucleotide sequences to be claimed in a single application. See Examination of Patent Applications Containing Nucleotide Sequences, 1192 O.G. 68 (November 19, 1996).

Although the MPEP deems that up to ten nucleotide sequences may be searched without restriction, it has recently been decided by the Director of Biotechnology at the USPTO that searching more than one sequence per application

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will place an undue burden upon the Examiner and the Office. For this reason, restriction to ONE SEQUENCE is being applied to all applications at this time.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Thaian N. Ton whose telephone number is (703) 305-1019. The examiner can normally be reached on Monday through Friday from 8:00 to 5:00 (Eastern Standard Time), with alternating Fridays off. Should the examiner be unavailable, inquiries should be directed to Deborah Reynolds, Supervisory Primary Examiner of Art Unit 1632, at (703) 305-4051. Any administrative or procedural questions should be directed to Patsy Zimmerman, Patent Analyst, at (703) 305-2758. Papers related to this application may be submitted to Group 1600 by facsimile transmission. Papers should be faxed to Group 1600 via the PTO Fax Center located in Crystal Mall 1. The faxing of such papers must conform with the notice published in the Official Gazette, 1096 OG 30 (November 15, 1989). The CM1 Fax Center number is (703)872-9306.

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DEBORAH CROUCH PRIMARY EXAMINER GROUP 1860 /430

TNT

Thaian N. Ton Patent Examiner Group 1632